FISCAL NOTE

HB 62

January 31, 2005

SUMMARY OF BILL: Requires informed consent documentation from all human subjects involved with any medical research. A copy of the written consent document signed by the subject or the subject's personal representative and an oral presentation of informed consent performed by a person knowledgeable about the details and risks of the research must be provided to the subject or the subject's personal representative. A non-biased third party witness must be present during the signing and oral presentation.

ESTIMATED FISCAL IMPACT:

State Expenditures - Net Impact - Not Significant - Health Related Boards

Assumption:

• The Department of Health would handle violations of this regulation in the same manner as all other disciplinary matters.

CERTIFICATION:

This is to duly certify that the information contained herein is true and correct to the best of my knowledge.

James W. White, Executive Director